

Hereditary benign telangiectasia?

To the Editor,

Takcı and colleagues¹ presented two 13 and 16 year old girls with telangiectasia in buccal mucosa and sun exposed skin areas under the title of “Hereditary Benign Telangiectasia” without family history, as the authors emphasized with supporting references.

When the ages of the patients are taken into consideration, I believe endoglin² gene mutation better be searched to separate these children’s telangiectasia from hereditary hemorrhagic telangiectasia (Osler-Rendu-Weber) syndrome which is usually seen in older people.

Şinasi Özsoylu, MD

Retired Professor of Pediatrics, Hematology and Hepatology

Honorary Fellow of American Academy of Pediatrics

Honorary Member of American Pediatric Society

Honorary Member of Turkish Academy of Sciences (TUBA)

Member of Islamic World Academy of Sciences (IAS)

E-mail: sinasiozsoylu@hotmail.com

PS: Unfortunately the pictures were not clear enough for evaluation.

REFERENCES

1. Takcı Z, Tekin Ö, Tezer A. Two cases of hereditary benign Telangiectasia in Turkey: sporadic occurrence with punctate telangiectasias surrounded by anemic halos. Turk J Pediatr 2015; 57: 94-97.
2. Perez-Belmonte LM, Gomez-Moyano E. Images in Clinical Medicine: Osler-Weber-Rendu syndrome. N Engl J Med 2015; 373: e15.